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APPLICATION FOR UNITED STATES LETTERS PATENT

S P E C I F I C A T I O N

TO ALL WHOM IT MAY CONCERN:

Be it known that I, **Lewis H. LAMBERT, Jr.**; a citizen of the United States of America, residing at the 45928 Omega Drive, Fremont, California 94539 have invented a new and useful **TREATMENT OF MYCOBACTERIAL DISEASES BY ADMINISTRATION OF BACTERICIDAL/PERMEABILITY-INCREASING PROTEIN PRODUCTS**, of which the following is a specification.

This is a continuation-in-part of co-pending U.S. Patent Application Serial No. 08/031,145, filed March 12, 1993.

5

The present invention relates to methods of treating a subject suffering from infection with *Mycobacteria* by administration of Bactericidal/Permeability-Increasing Protein (BPI) protein products.

Mycobacterium is a non-motile, acid-fast, aerobic, genus of bacteria known to cause grave human and animal diseases, such as tuberculosis and leprosy. Infections caused by *M. avium* are the most common form of disseminated bacterial disease in AIDS patients. Orme, *et al.*, *Infect. and Immun.*, 61(1):338-342 (1993).

The administration of conventional antibiotics to treat *Mycobacterial* infection is known in the art and has achieved varying success depending on the susceptibility of the bacterial strain, the efficacy and toxicity of the antibiotic(s) employed, the duration of treatment, and numerous other factors. Antimicrobials that have been employed alone or in combination to treat *Mycobacterial* infections, including those caused by *M. tuberculosis* include isoniazid, rifampin, ethambutol, *p*-aminosalicylic acid, pyrazinamide, streptomycin, capreomycin, cycloserine, ethionamide, kanamycin, amikacin, amithiozone, rifabutin, clofazimine, arithromycin, clarithromycin, ciprofloxacin and ofloxacin. McClatchy, *Antimycobacterial Drugs: Mechanisms of Action, Drug Resistance, Susceptibility Testing, and Assays of Activity in Biological Fluids*, pp. 134-197, In Antibiotics in Laboratory Medicine, 3rd ed., V. Lorian, ed. The Williams & Wilkins Co., Baltimore (1991). As many *Mycobacterial* strains are drug resistant, serious obstacles

exist for control and successful treatment of tuberculosis and other *Mycobacterial* diseases. *Id.*

A variety of factors have made treatment of individuals afflicted with *Mycobacterial* diseases problematic. First, *Mycobacteria* possess a very hydrophobic cell wall that affords protection against the host's immune system.

As *Mycobacterial* infections tend to be chronic, the pathologies of these organisms are generally due to host response. Also, many *Mycobacterial* strains are drug-resistant. These and other factors make the development of novel, effective methods for treating *Mycobacterial* diseases highly desirable.

Mycobacteria are readily distinguished from gram-negative and gram-positive bacteria by acid fast staining due to significant differences in cell wall structure. Gram-negative bacteria are characterized by a cell wall composed of a thin layer of peptidoglycan covered by an outer membrane of lipoprotein and lipopolysaccharide (LPS), whereas gram-positive bacteria have a cell wall with a thicker layer of peptidoglycan with attached teichoic acids, but no LPS. The *Mycobacterial* cell wall is rich in fatty acids, including a major constituent, lipoarabinomannan (LAM), which is widely distributed within the cell wall of *Mycobacterium* species. LAM has been purified from both *M. leprae* and *M. tuberculosis*. Hunter *et al.*, *J Biol. Chem.*, 261:12345-12351 (1986). LAM is a serologically active mannose containing phosphorylated lipopolysaccharide that may be membrane associated.

The complex physiological effects of LAM appear to be concentration, time, and source-dependent. For example, Chatterjee *et al.*, *Infect. and Immun.*, 60(3):1249-1253 (1992), reported that, in the first 24 hours following exposure, LAM from an avirulent strain of tuberculosis was 100-fold more potent at stimulating TNF secretion in mouse macrophages than LAM from a virulent strain. LAM concentrations of 0.01-10 $\mu\text{g/ml}$ for the avirulent strain and 0.01-100 $\mu\text{g/ml}$ for the virulent strain were tested, and increased LAM concentration was associated with increased TNF production with LAM from both species.

Macrophage-inhibitory effects of LAM have also been described in the art. LAM purified from both *M. leprae* and *M. tuberculosis* has been reported to be a potent *in vitro* inhibitor of T-cell lymphokine activation of mouse macrophages. Sibley *et al.*, *Infection and Immunity*, 56(5):1232-1236 (1988). Because the principle efferent role of the macrophage in acquired resistance to intracellular pathogens requires activation by T-cell lymphokines, notably gamma-interferon (IFN- γ), macrophages whose activation-response is inhibited are severely compromised in their capacity for both enhanced microbicidal and tumoricidal activities.

In another study, Sibley *et al.*, *Clin. Exp. Immunol.*, 80(1):141-148 (1990), reported that pretreatment of mouse macrophages with 50 to 100 μ g/ml LAM blocked macrophage activation by IFN- γ , but pretreatment with 10 μ g/ml LAM did not affect macrophage activation. Thus, it is believed that low concentrations of LAM stimulate cytokine production, at least initially. However, higher concentrations of LAM (50-100 μ g/ml or more) appear to block rather than promote macrophage function. Thus, the production of either too much or too little cytokine at different stages of *Mycobacterial* disease may contribute to *Mycobacterial* pathogenesis. New methods for blocking the above-characterized physiological effects of LAM molecules are a highly desirable goal in the treatment of subjects that are or that have been infected with *Mycobacteria*. For the same reasons, new methods by which fluids containing LAM can be decontaminated prior to administration into a subject are also desirable. Neutralization of even small amounts of LAM is desirable, because small amounts of LAM may have the physiological effect of stimulating cytokine production.

Of interest to the background of the invention are the disclosures of PCT/US88/00510, (WO 88/06038) published August 25, 1988, indicating that certain poloxypropylene/polyoxyethylene nonionic surface-active block copolymers can be used with or without conventional antibiotics to treat infection with *Mycobacterium*. This reference cites studies suggesting that the

effects of other nonionic surfactants on tuberculosis are most likely due to modification of surface lipids of *Mycobacteria*, and not to direct bactericidal effects on *Mycobacteria*. See e.g. Cornforth *et al.*, *Nature*, 168:150-153 (1951).

5 Bactericidal/permeability-increasing protein (BPI) is a protein isolated from the granules of mammalian polymorphonuclear neutrophils (PMN), which are blood cells essential in the defense against invading microorganisms. Human BPI protein has been isolated from PMN's by acid extraction combined with either ion exchange chromatography Elsbach, *J.*
10 *Biol. Chem.*, 254:11000 (1979) or *E. coli* affinity chromatography, Weiss, *et al.*, *Blood*, 69: 652 (1987), and has potent bactericidal activity against a broad spectrum of gram-negative bacteria. The molecular weight of human BPI is approximately 55,000 Daltons (55 kD). The amino acid sequence of the entire human BPI protein, as well as the DNA encoding the protein, have been
15 elucidated in Figure 1 of Gray, *et al.*, *J. Biol. Chem.*, 264: 9505 (1989), incorporated herein by reference.

BPI has been shown to be a potent bactericidal agent active against a broad range of gram-negative bacterial species. The cytotoxic effect of BPI was originally established to be highly specific to sensitive gram-
20 negative species, with no toxicity being noted for other non-acid fast, gram-positive bacteria or for eukaryotic cells. The precise mechanism by which BPI kills bacteria is as yet unknown, but it is known that BPI must first attach to the surface of susceptible gram-negative bacteria. It is thought that this initial binding of BPI to the bacteria involves electrostatic interactions between the
25 basic BPI protein and negatively charged sites on lipopolysaccharides (LPS). LPS has been referred to as endotoxin because of the potent inflammatory response that it stimulates. LPS induces the release of mediators by host inflammatory cells which may ultimately result in irreversible endotoxic shock. BPI binds to Lipid A, the most toxic and most biologically active component
30 of LPS.

In susceptible bacteria, it is thought that BPI binding disrupts LPS structure, leads to an activation of bacterial enzymes that degrade phospholipids and peptidoglycans, alters the permeability of the cell's outer membrane, and ultimately causes cell death by an as yet unknown mechanism.

5 BPI is also capable of neutralizing the endotoxic properties of LPS to which it binds. Because of its gram-negative bactericidal properties and its ability to neutralize LPS, BPI can be utilized for the treatment of mammals suffering from diseases caused by gram-negative bacteria, such as bacteremia or sepsis.

An approximately 25 kD proteolytic fragment corresponding to
10 the amino-terminal portion of human BPI holoprotein possesses the antibacterial efficacy of the naturally-derived 55 kD human holoprotein. In contrast to the amino-terminal portion the carboxy-terminal region of the isolated human BPI protein displays only slightly detectable anti-bacterial activity. Ooi, *et al.*, *J. Exp. Med.*, 174:649 (1991). A BPI amino-terminal
15 fragment, expressed from a construct encoding approximately the first 199 amino acid residues of the human BPI holoprotein, has been produced by recombinant means as a 23 kD protein referred to as "rBPI₂₃". Gazzano-Santoro *et al.*, *Infect. Immun.* 60: 4754-4761 (1992).

While BPI protein products are effective for treatment of
20 conditions associated with gram-negative bacterial infection, there continues to exist a need in the art for products and methods for treatment of other bacterial infections such as infection with *Mycobacteria*.

SUMMARY OF THE INVENTION

25 The present invention provides methods of treating a subject suffering from infection with *Mycobacteria* by administration of a composition comprising a BPI protein product. Therapeutic compositions according to the invention may be administered orally, systemically (such as by intravenous, intramuscular or other injection), or as an aerosol. *Mycobacterial* disease
30 states subject to treatment according to the invention include tuberculosis,

which can be caused by infection with *M. tuberculosis*, leprosy, which can be caused by infection with *M. leprae*, and diseases caused by *M. avium* and other *Mycobacteria* species. According to preferred methods, anti-Mycobacterial antibiotics such as previously identified and/or surfactants may be administered
5 in combination with the BPI protein product to subjects suffering from infection with *Mycobacteria*.

According to another aspect of the present invention, compositions comprising a BPI protein product are administered to neutralize LAM's physiological effects on a host. For example, methods are provided
10 for neutralizing the effect of low concentrations of LAM capable of stimulating cytokine production in a host. Methods are also provided for neutralizing the inhibitory effect that higher concentrations of Mycobacterial LAM (i.e. 100 $\mu\text{g/ml}$ or more) have upon the interferon-mediated activation of macrophages. Specifically, a BPI protein product may be administered to an
15 immunosuppressed subject failing to respond to microbes or tumor cells due to LAM-induced insensitivity of macrophages to activation by T-cell lymphokines.

According to a further aspect of the present invention, a BPI protein product is employed in methods for decontaminating a fluid containing
20 LAM prior to administration of the fluid into a subject. Such decontamination methods of the invention involve contacting the fluid with the BPI protein product prior to administration, under conditions such that LAM forms a complex with the BPI protein product which can be removed from the fluid. Fluids subject to decontamination by the methods of this invention include, but
25 are not limited to, blood, plasma, blood serum, bone marrow, isotonic solutions, pharmaceutical agents, and cell culture agents.

A further aspect of this invention relates to the use of a composition comprising a BPI protein product for the manufacture of a medicament for the therapeutic application of treating any of the
30 aforementioned conditions or infections from which a subject might suffer.

Numerous additional aspects and advantages of the invention will become apparent to those skilled in the art upon considering the following detailed description of the invention, which describes presently preferred embodiments thereof.

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BRIEF DESCRIPTION OF THE FIGURES

Fig. 1 graphically depicts the results of an assay of BPI protein product binding to *E. coli* J5 Lipid A and *M. tuberculosis* and Fig. 2 graphically represents the results of test to assess the ability of a BPI protein product to inhibit mycobacterial induced TNF production in whole blood.

10

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to the discovery that a composition comprising a BPI protein product can be administered for effective treatment of a subject suffering from infection with *Mycobacteria*. In particular, the invention provides methods for treatment of leprosy and tuberculosis, grave diseases caused by the species *M. leprae* and *M. tuberculosis*, respectively. It is contemplated that the methods described herein may be used to treat infection with other *Mycobacterial* species, most notably *M. avium* and *M. intracellulare* (collectively known as "MAC"), but also *M. marinum*, *M. fortuitum*, *M. chelonae*, *M. smegmatis*, *M. kansasii*, *M. bovis*, *M. hominis*, *M. gordonae* and other pathogenic or opportunistic species. Beneficial effects of treatment with BPI protein products are expected to result from binding of the products to LAM and disruption of the bacterial cell wall components (with or without direct killing of the bacteria) in manner similar to that resulting from treatment of gram-negative disease states.

25

As used herein, "BPI protein product" includes naturally and recombinantly produced BPI protein; natural, synthetic, and recombinant biologically active polypeptide fragments of BPI protein; biologically active polypeptide variants of BPI protein or fragments thereof, including hybrid

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fusion proteins and dimers; and biologically active polypeptide analogs of BPI protein or fragments or variants thereof, including cysteine-substituted analogs. The BPI protein products administered according to this invention may be generated and/or isolated by any means known in the art. U.S. Patent No. 5,198,541, the disclosure of which is hereby incorporated by reference, discloses recombinant genes encoding and methods for expression of BPI proteins including recombinant BPI holoprotein, referred to herein as rBPI₅₀ and recombinant fragments of BPI. Co-owned, copending U.S. Patent Application Ser. No. 07/885,501 and a continuation-in-part thereof, U.S. Patent Application Ser. No. 08/072,063 filed May 19, 1993 which are hereby incorporated by reference, disclose novel methods for the purification of recombinant BPI protein products expressed in and secreted from genetically transformed mammalian host cells in culture and discloses how one may produce large quantities of recombinant BPI products suitable for incorporation into stable, homogeneous pharmaceutical preparations.

Biologically active fragments of BPI (BPI fragments) include biologically active molecules that have the same amino acid sequence as a natural human BPI holoprotein, except that the fragment molecule lacks amino-terminal amino acids, internal amino acids, and/or carboxy-terminal amino acids of the holoprotein. Nonlimiting examples of such fragments include an N-terminal fragment of natural human BPI of approximately 25 kD, described in Ooi et al., *J. Exp. Med.*, 174:649 (1991), and the recombinant expression product of DNA encoding N-terminal amino acids from residue 1 to about residue 200, including from about residue 1 to about residue 193 or 199 of natural human BPI, described in Gazzano-Santoro et al., *Infect. Immun.* 60:4754-4761 (1992), and referred to as rBPI₂₃. In that publication, an expression vector was used as a source of DNA encoding a recombinant expression product (rBPI₂₃) having the 31-residue signal sequence and the first 199 amino acids of the N-terminus of the mature human BPI, as set out in Figure 1 of Gray et al., *supra*, except that valine at position 151 is specified

by GTG rather than GTC and residue 185 is glutamic acid (specified by GAG) rather than lysine (specified by AAG). Recombinant holoprotein (rBPI) has also been produced having the sequence (SEQ. ID NOS: 1 and 2) set out in Figure 1 of Gray et al., *supra*, with the exceptions noted for rBPI₂₃ and with the exception that residue 417 is alanine (specified by GCT) rather than valine (specified by GTT). Other examples include dimeric BPI forms as described in co-owned and co-pending U.S. Patent Application Serial No. 08/212,132, filed March 11, 1994, the disclosure of which is hereby incorporated by reference.

Biologically active variants of BPI (BPI variants) include but are not limited to recombinant hybrid fusion proteins, comprising BPI holoprotein or a biologically active fragment thereof and at least a portion of at least one other polypeptide, and dimeric forms of BPI variants. Examples of such hybrid fusion proteins and dimeric forms are described by Theofan et al. in co-owned, copending U.S. Patent Application Serial No. 07/885,911, and a continuation-in-part application thereof U.S. Patent Application Serial No. 08/064,693 filed May 19, 1993 which are incorporated herein by reference in their entirety and include hybrid fusion proteins comprising, at the amino-terminal end, a BPI protein or a biologically active fragment thereof and, at the carboxy-terminal end, at least one constant domain of an immunoglobulin heavy chain or allelic variant thereof.

Biologically active analogs of BPI (BPI analogs) include but are not limited to BPI protein products wherein one or more amino acid residue has been replaced by a different amino acid. For example, co-owned, copending U.S. Patent Application Ser. No. 08/013,801 (Theofan et al., "Stable Bactericidal/Permeability-Increasing Protein Products and Pharmaceutical Compositions Containing the Same," filed February 2, 1993), the disclosure of which is incorporated herein by reference, discloses polypeptide analogs of BPI and BPI fragments wherein a cysteine residue is replaced by a different amino acid. A preferred BPI protein product described

by this application is the expression product of DNA encoding from amino acid 1 to approximately 193 or 199 of the N-terminal amino acids of BPI holoprotein, but wherein the cysteine at residue number 132 is substituted with alanine and is designated rBPI₂₁Δcys or rBPI₂₁.

5 Other BPI protein products useful according to the methods of the invention are peptides derived from or based on BPI produced by recombinant or synthetic means (BPI-derived peptides), such as those described in co-owned and copending U.S. Patent Application Serial No. 08/209,762, filed March 11, 1994, which is a continuation-in-part of U.S. Patent
10 Application Serial No. 08/183,222, filed January 14, 1994, which is a continuation-in-part of U.S. Patent Application Ser. No. 08/093,202 filed July 15, 1993), which is a continuation-in-part of U.S. Patent Application Ser. No. 08/030,644 filed March 12, 1993, the disclosures of which are hereby incorporated by reference. Other useful BPI protein products include peptides
15 based on or derived from BPI which are described in co-owned and co-pending U.S. Patent Application Serial No. 08/274,299 filed July 11, 1994, by Horwitz *et al.* and U.S. Patent Application Serial No. 08/273,540, filed July 11, 1994, by Little *et al.*

 Presently preferred BPI protein products include recombinantly-produced N-terminal fragments of BPI, especially those having a molecular
20 weight of approximately between 21 to 25 kD such as rBPI₂₁ or rBPI₂₃, dimeric forms of these N-terminal fragments. Additionally, preferred BPI protein products include rBPI₅₀ and BPI-derived peptides.

 The administration of BPI protein products is preferably
25 accomplished with a pharmaceutical composition comprising a BPI protein product and a pharmaceutically acceptable diluent, adjuvant, or carrier. The BPI protein product may be administered without or in conjunction with known surfactants, other chemotherapeutic agents. A preferred pharmaceutical composition containing BPI protein products comprises the BPI protein product
30 at a concentration of 1 mg/ml in citrate buffered saline (5 or 20 mM citrate,

150 mM NaCl, pH 5.0) comprising 0.1% by weight of poloxamer 188 (Pluronic F-68, BASF Wyandotte, Parsippany, NJ) and 0.002% by weight of polysorbate 80 (Tween 80, ICI Americas Inc., Wilmington, DE). Another preferred pharmaceutical composition containing BPI protein products
5 comprises the BPI protein product at a concentration of 2 mg/ml in 5 mM citrate, 150 mM NaCl, 0.2% poloxamer 188 and 0.002% polysorbate 80. Such preferred combinations are described in co-owned, co-pending, U.S. Patent Application Ser. No. 08/190,869 filed February 2, 1994 (McGregor et al., "Improved Pharmaceutical Compositions"), and U.S. Patent Application
10 Ser. No. 08/012,360 filed February 2, 1993 (McGregor et al., "Improved Pharmaceutical Composition"), the disclosures of which are incorporated herein by reference.

The BPI protein product can be administered by any known method, such as orally, systemically (such as by intravenous, intramuscular or
15 other injection), or as an aerosol. Medicaments can be prepared for oral administration or by injection or other parenteral methods and preferably include conventional pharmaceutically acceptable carriers and adjuvants as would be known to those of skill in the art. The medicaments may be in the form of a unit dose in solid, semi-solid and liquid dosage forms such as
20 tablets, pills, powders, liquid solutions or suspensions, and injectable and infusible solutions. Effective dosage ranges from about 100 μ g/kg to about 10 mg/kg of body weight are contemplated. Intravenous administration is a preferred method for treatment of leprosy.

It is contemplated that aerosol administration to the lungs will be
25 a preferred method for treating other *Mycobacterial* infections, such as tuberculosis. Such aerosol formulations would be manufactured by means that are known in the art, and administered by metered-dose inhaler, updraft nebulization, or other means known in the art.

An aspect of the present invention is to provide methods of
30 treating a subject suffering from any of the physiological effects of

Mycobacterial LAM. As described above, the physiological effects of LAM depend on a number of factors, including the source and concentration of the LAM, and the length of time to which host cells are exposed to LAM. Example 3, infra, demonstrates that 20 - 100 $\mu\text{g/ml}$ of nonviable, desiccated

5 *M. tuberculosis* added to whole blood will stimulate TNF production by the monocytes in the blood. Other studies described above have shown that 50-100 $\mu\text{g/ml}$ of LAM will down-regulate macrophage functions and expression (TNF, and the like) and prevent macrophage activation, said methods comprise administering a BPI protein product to the subject. Methods are provided for

10 treating a subject suffering from the effects of increased cytokine production caused by the physiological presence of LAM. Methods are also provided for treating a subject suffering from LAM-induced inhibition of macrophage activation, and the effects thereof. Methods and formulations by which a BPI protein product may be administered, including preferred methods and

15 formulations, are the same as those set forth above for the treatment of *Mycobacterial* infection.

Because of the harmful physiological effects that *Mycobacterial* LAM can have on a subject, even in the absence of viable *Mycobacteria*, methods are provided in the present invention by which a fluid containing

20 LAM may be decontaminated prior to administration of the fluid into a subject. Such methods comprise contacting the fluid with a BPI protein product prior to administration, under conditions such that LAM forms a complex with the BPI protein product, thereby decontaminating the fluid. By way of nonlimiting examples, such methods may be applied to fluids such as blood, plasma, blood

25 serum, bone marrow, isotonic solutions, pharmaceutical agents, or cell culture reagents.

BPI protein product is thought to interact with a variety of host defense elements present in whole blood or serum, including complement and LBP, and other cells and components of the immune system. Such interactions

30 might result in potentiating and/or synergizing the anti-microbial activities.

Because of these interactions, BPI protein products are expected to exert even greater activity *in vivo* than *in vitro*. Thus, while *in vitro* tests are predictive of *in vivo* utility, absence of activity *in vitro* does not necessarily indicate absence of activity *in vivo*. For example, BPI has been observed to display a greater bactericidal effect on certain gram-negative bacteria in whole blood or plasma assays than in assays using conventional media. [Weiss et al., *J. Clin. Invest.* 90:1122-1130 (1992)]. This may be because conventional *in vitro* systems lack the blood elements that facilitate or potentiate BPI's function *in vivo*, or because conventional media designed to maximize bacterial growth contain higher than physiological concentrations of magnesium and calcium, inhibitors of BPI protein product antibacterial activity.

Therapeutic effectiveness is based on a successful clinical outcome, and does not require that an anti-mycobacterial agent or agents kill 100% of the organism involved in the infection. Frequently, reducing organism load by one log (factor of 10) permits the host's own defenses to control the infection. In addition, augmenting an early anti-mycobacterial effect can be particularly important in addition to any long-term anti-mycobacterial effect. These early events are a significant and critical part of therapeutic success, because they allow time for host defense mechanisms to activate.

It is also contemplated that the BPI protein product be administered with other products that potentiate the anti-mycobacterial activity of BPI protein products. For example, serum complement potentiates the gram-negative bactericidal activity of BPI protein products; the combination of BPI protein product and serum complement provides synergistic bactericidal/growth inhibitory effects. See, e.g., Ooi *et al. J. Biol. Chem.*, 265: 15956 (1990) and Levy *et al. J. Biol. Chem.*, 268: 6038-6083 (1993) which address naturally-occurring 15 kD proteins potentiating BPI antibacterial activity. See also co-owned, co-pending U.S. Patent Application Serial No. 08/093,201 filed July 14, 1993, and continuation-in-part, U.S. Patent

Application Serial No. 08/274,303 filed July 11, 1994 which describes methods for potentiating gram-negative bactericidal activity of BPI protein products by administering lipopolysaccharide binding protein (LBP) and LBP protein products. The disclosures of these applications are incorporated by
5 reference herein. LBP protein derivatives and derivative hybrids which lack CD-14 immunostimulatory properties are described in co-owned, co-pending U.S. Patent Application Serial No. 08/261,660, filed June 17, 1994 as a continuation-in-part of U.S. Patent Application Serial No. 08/079,510, filed June 17, 1993, the disclosures of which are incorporated by reference herein.

10 An aspect of this invention includes the use of a composition comprising a BPI protein product for the manufacture of a medicament for the therapeutic application of treating any of the aforementioned conditions or diseases from which a subject suffers. The medicament may include, in addition to a BPI protein product, other chemotherapeutic agents such as
15 known anti-mycobacterial antibiotics or surfactants. The medicament may additionally or alternatively include one or more additional pharmaceutically acceptable components, such as diluents, adjuvants, or carriers.

An aspect of the present invention is the ability to provide more effective treatment of Mycobacterial infection by virtue of the synergistic
20 increase in or potentiation of the anti-bacterial activities of an anti-Mycobacterial antibiotic or BPI protein product. As previously noted, anti-Mycobacterial antibiotic therapy currently involves administration of one or more (and frequently three or more) antibiotics such as isoniazid, rifampin, ethambutol, *p*-aminosalicylic acid, pyrazinamide, streptomycin, capreomycin,
25 cycloserine, ethionamide, kanamycin, amikacin, amithiozone, rifabutin, clofazimine, arithromycin, clarithromycin, ciprofloxacin and ofloxacin. Unlike some therapeutic agents, BPI protein product is easily administered and produces no inflammatory reaction. An aspect of the present invention is the ability to treat Mycobacterial organisms that are normally resistant to one or
30 more antibiotics. A further aspect is the ability to use lower concentrations of

relatively toxic or expensive antibiotics such as rifampin. Because the use of some antibiotics is limited by their systemic toxicity or prohibitive cost, lowering the concentration of antibiotic required for therapeutic effectiveness reduces toxicity and/or cost of treatment, and thus allows wider use of the antibiotic. The present invention may also provide quality of life benefits due to, e.g., decreased duration of therapy, reduced stay in intensive care units or overall in the hospital, with the concomitant reduced risk of serious nosocomial (hospital-acquired) infections.

The invention further provides pharmaceutical compositions for treatment of Mycobacterial infection and the sequelae thereof comprising the combination of a BPI protein product and an antibiotic which is present in an amount effective to have synergistic or potentiating bactericidal/bacteriostatic properties, including increased susceptibility or reversal of resistance. The pharmaceutical composition can comprise a pharmaceutically-acceptable diluent, adjuvant or carrier.

Methods of the present invention are more fully illustrated by the nonlimiting examples which follow. Example 1 address BPI protein products binding to a species of *Mycobacterium*, *M. tuberculosis*. Example 2 address prospective use of BPI protein products in binding purified LAM of *Mycobacteria*. Examples 3 and 4 describe attempts to reverse *Mycobacteria*-induced cytokine production in whole human blood. Example 5 addresses use of BPI protein products in combination with anti-mycobacterial antibiotics to inhibit *M. tuberculosis* growth. Remaining Examples 6-13 address prospective *in vitro* and *in vivo* use of BPI protein products according to methods of this invention. The models described in those examples and/or other models known in the art are used to predict the efficacy and the optimal BPI protein product formulations of the methods of invention.

EXAMPLE 1

An enzyme linked immunosorbent assay (ELISA) was conducted to determine binding of a BPI protein product to *M. tuberculosis*. Specifically, non-viable, desiccated *M. tuberculosis* H37 RA (Difco, Detroit MI) was
5 suspended in DPBS (25 μ g/ml) and used to coat microtiter wells overnight at 37°C. Wells were also coated with either 25 μ g/ml Lipid A (*E.coli* J5 mutant, RIBI, Hamilton MT) or 500 μ l DPBS to demonstrate the functionality and specificity of rBPI₂₃. After washing (3x with DPBS + 0.05% Tween 20), the plates were blocked for 1hr. at room temperature with 200 μ l/well of DPBS +
10 1% non-fat milk. After washing as above, 50 μ l solutions of either various concentrations of rBPI₂₃ (in DPBS containing 0.05% Tween 20) or DPBS (negative control) were added to the wells, which were then incubated for 1 hr. at 37°C. The wells were again washed as above, and the amount of rBPI₂₃ bound to the wells was determined using an anti-rBPI₂₃ mouse monoclonal
15 antibody (designated α BPI MAb-2-4) and an enzyme conjugated anti-murine IgG antibody (HRP-Ab, Zymed #61-0120, San Francisco, CA). To each well 100 μ l of α BPI MAb-2-4 was added (100 ng/ml in DPBS + 0.05% Tween 20), and the plates were incubated 1 hr. at 37°C. After washing as above, 100 μ l of HRP-Ab was added (1:1000 in DPBS + 0.05% Tween 20) to each well
20 and the plates were again incubated 1 hr. at 37°C. After washing the plates as above, 100 μ l substrate in 0.1M citrate plus 1:50 ABTS (20 mg/ml stock) and 1:1000 H₂O₂ was added to each well. The plates were incubated 10-30 min. at room temperature, and absorbance readings were taken at 405 nm (OD 405).

The results of the experiment are represented graphically in
25 Figure 1, which depicts the ability of varying concentrations rBPI₂₃ to bind to J5 Lipid A (filled triangles); to *M. tuberculosis* (open squares); and to the no antigen-free control (filled circles). The abscissa of each measurement represents the concentration of rBPI₂₃, and the ordinate represents the average OD 405 measurements from four trials. Error bars reflect the variation in OD
30 405 readings for each data point.

This experiment demonstrated that rBPI₂₃ binds specifically to non-viable desiccated *M.tuberculosis*. The functionality of the rBPI₂₃ used in these experiments was confirmed by the results of the Lipid A (positive control) binding assay, and the specificity of the experiments was confirmed by the lack of binding to the negative control samples.

EXAMPLE 2

In this example, an ELISA Assay is conducted to determine binding of a BPI protein product to the lipoarabinomannan portion of *Mycobacteria*. The binding activity of BPI protein product (e.g., rBPI₂₃) to LAM is demonstrated as described in the previous example, except LAM purified from a species of *Mycobacterium*, (e.g., *M.tuberculosis* or *M.leprae*) is substituted for the nonviable *M.tuberculosis* used to coat the ELISA plates in that example. Purified LAM is isolated as described by Hunter *et al.*, *J. Biol. Chem.*, 261:12345-12351 (1986). Specific binding of biologically active BPI protein product is demonstrated by comparison of the OD 405 readings from the LAM coated wells with positive and negative controls.

EXAMPLE 3

The following experiment was conducted to determine the effect of a BPI protein product, rBPI₂₃, on *Mycobacteria*-induced cytokine production in whole human blood. Whole human blood from healthy volunteers was collected into Vacutainer tubes (ACD, Beckton Dickinson, Rutherford, NJ). Aliquots of blood (225 μ l) were mixed with either rBPI₂₃ (10 μ g/ml final) or the protein thaumatin (10 μ g/ml final in 5 ml) as a negative control. RPMI medium (20 μ l) was added to each sample. Varying dilutions (0 - 8 ng/ml) of either *E. coli* O113 LPS (Ribi, Hamilton MT) or of non-viable, desiccated *M. tuberculosis* H37 RA (0 - 100 μ g/ml) (Difco, Detroit MI) were added to the samples, which were then incubated at 37°C for 6 hours. The reactions were stopped by the addition of 750 μ l of RPMI medium, the samples were

centrifuged at 500 g for 7 min, and stored at -20°C until analyzed. The supernatant was assayed for cytokine (TNF) levels based on a standard curve, according to the manufacturers' recommendation (Biokine ELISA test, T Cell Sciences, Cambridge, MA).

5 The assay results revealed that rBPI₂₃ at 10 µg/ml had no inhibitory effect on *M. tuberculosis*-induced TNF release at the concentration (20 - 100 µg/ml) of *M. tuberculosis* added to the blood samples. The same concentration of rBPI₂₃ eliminated LPS-induced TNF release at the LPS concentrations tested (2 - 8 ng/ml). The lack of inhibitory effect on cytokine
10 induction by *M. tuberculosis* may be the result of use of sub-optimal dosage levels. Alternatively, some component of the *Mycobacterial* cell wall other than the LAM bound by rBPI₂₃ may be responsible for inducing cytokine production at the *Mycobacterium* concentrations tested.

EXAMPLE 4

In this example, multiple additional assays were conducted to assess the inhibitory effect of BPI protein products on mycobacterial (*M. tuberculosis* or *M. smegmatis*) induced production of tumor necrosis factor (TNF) by monocytes/macrophages present in whole human blood. Briefly summarized, live or heat killed mycobacteria at varying concentrations was added to whole blood and incubated with either a fixed amount BPI protein product or acetate buffer negative control solution. After incubation, the content of TNF present was assessed by standard means. The TNF content of BPI protein product treated samples was then compared to the TNF content of buffer control samples to determine the relative inhibitory effect of the BPI protein product tested. Whole blood samples were obtained from healthy human volunteers and aliquoted as in Example 3. To each tube containing 225 μ l of whole blood was added from 0 to 1×10^7 live or heat killed mycobacteria. Depending on whether one or two BPI protein products were to be tested, four or six tubes were prepared at each concentration of bacteria.

after which the tubes were incubated at 37°C for 15 minutes. To two of the tubes at each bacterial concentration was added either a selected BPI protein product at a final concentration of 4 µg/ml or acetate buffer (as a negative control), after which is tubes were further incubated at 37°C for 5 to 6 hours.

- 5 Thereafter 750 µl of RPMI 1640 was added to each sample and the samples were centrifuged at 17,000 rpm (500 g?) for 6 minutes. Supernatants were stored at -70°C until thawed immediately prior to testing for TNF content using the Biokine ELISA test kit as in Example 3.

- 10 In a first series of assays, heat-killed *M. tuberculosis* (strain H37Ra) was employed as the TNF stimulating organism. In a first test on whole blood, no substantial increased in TNF levels was observed until bacteria were added at a concentration of 1×10^6 organisms and the presence of rBPI₂₃ in the samples resulted in an approximately 70% inhibition of TNF production. In a second test involving two separate whole blood assays,
- 15 inhibitory effects of both rBPI₂₃ and BPI holoprotein were assessed. Substantial increases in TNF concentration over basal levels (no microorganisms added) were observed commencing at microorganism concentrations of 3×10^5 up through 1×10^7 . Overall, inhibition of TNF production by 20% or greater was observed when rBPI₂₃ was added at all such
- 20 organism levels. Lesser degrees of inhibition were noted for the BPI holoprotein (with no inhibition at all noted in one duplicate test at the highest concentration of organisms). The lesser effects of the holoprotein in these assays are likely attributable to the lower molar concentration employed. A third *M. tuberculosis* test was performed on whole blood drawn from four
- 25 different volunteers, using rBPI₂₃ as the test compound. Expectedly, the level of inhibition of TNF formation by the uniform dose of BPI protein product varied from subject to subject. With the exception of one subject's blood samples (wherein inhibition was observed only at intermediate microorganism concentration of 1×10^6 and not at all at concentrations of 1×10^7), the BPI
- 30 protein product provided for at least about 10% and up to about 50% TNF

inhibition, with higher inhibitory levels being observed at higher microorganism concentrations. A fourth test involving *M. tuberculosis* was carried out using rBPI₂₃ at 4 µg/ml and 8 µg/ml concentrations. The BPI protein product was again observed to inhibit TNF at the 4 µg/ml level, with the

5 greatest effects being observed at microorganism concentrations of 3×10^6 . Doubling the concentration of test compound to 8 µg/ml did not enhance, and in fact somewhat diminished, inhibitory effects observed.

In a second series of assays, heat killed *M. smegmatis* was employed to stimulate TNF production in whole blood. In a first test, whole

10 blood from eight different patients was employed and was subjected to contact with concentrations of 0, 0.5×10^5 , 1×10^5 , 1×10^6 and 1×10^7 organisms. Figure 2 provides a graphic representation of the sum of the results observed and indicated that the rBPI₂₃ product tested at 4 µg/ml was an effective inhibitor of TNF production at all bacterial concentrations. In a second test

15 involving blood from two different subjects, TNF production inhibitory effects of rBPI₂₃ were assessed for *M. smegmatis*, *E. coli* and *S. aureus*. Expectedly, significant TNF inhibitory effects were observed in the *E. coli* treated blood, with the greatest present with microorganism concentrations of 1×10^5 and the lesser effects at higher concentrations of organisms. Similarly, no substantial

20 TNF inhibitory effects were observed for the BPI protein product in the *S. aureus* assay. Variable results were seen in the *M. smegmatis* assay; pronounced inhibitory effects were observed in one subject's blood at the 1×10^7 concentration of organisms, while inhibition was observed in the other blood sample only at the 1×10^6 microorganism concentration.

25 The above assay results demonstrate *in vitro* effectiveness of BPI protein products in inhibiting induction of tumor necrosis factor by mycobacterial species and are predictive of *in vivo* efficacy in human patients.

EXAMPLE 5

In this example, rBPI₂₃ at varying concentrations was assessed for its growth inhibitory effect on *M. tuberculosis* treated with varying concentrations of the anti-Mycobacterial antibiotics isoniazid (INH) and rifampin (RMP). Briefly summarized, pure cultures of *M. tuberculosis* (MTB) were incubated for 24 hours with varying concentrations of rBPI₂₃ and antibiotic. Cultures were added to Bactec[®] bottles (Johnston Laboratories, Cockeysville, Maryland) containing ¹⁴C labeled nutrients and daily "growth index" values were determined accordingly to the supplier's instructions on the basis of ¹⁴CO₂ evolved from the medium. In separate assays, concentrations of rBPI₂₃ of 0, 3.9, 15.6, 62.5, 250 and 1000 µg/ml were combined with INH at levels of 0, 0.006, 0.012, 0.025 and 0.05 µg/ml or RMP levels of 0, 0.12, 0.25, 0.5 and 1.0 µg/ml. Growth index values were assessed daily starting the second day after inoculation into the vials through to the eighteenth day. In the INH assay, no BPI protein product was added, growth index values characteristically gradually increased over time and as a function of the dosage of antibiotic employed (increases generally began earlier and rose more steeply at lower doses than at higher doses). Addition of rBPI₂₃ had variable effects in enhancing or diminishing antibiotic effects on growth index values, depending on the concentration employed. An intermediate dose (62.5 µg/ml) of BPI protein product consistently tended to reduce growth index values at all doses of INH tested and thus operated to enhance INH growth inhibitory effects. Similar but less pronounced enhancement effects were observed for the 15.6 µg/ml rBPI₂₃ dose. Lower (0 and 3.9 µg/ml) and higher (250 and 1000 µg/ml) doses of the BPI protein product generally diminished the antibiotic effects of INH, with the highest rBPI₂₃ dose invariably functioning to increasing "swamp out" INH effects on growth index toward the middle of the test period. At later times in the test period, however, the highest doses of BPI protein product appeared to suppress and actually reverse the above-noted characteristic increases in growth index over time. In combinative assay with

RMP, rBPI₂₃ had no discernible enhancing effect at the highest doses of the antibiotic 0.5 and 1.0 $\mu\text{g/ml}$ where there was essentially no increase in growth index throughout the entire test period. At lesser concentrations of RMP, there tended to be a dose-dependent enhancement effect of the BPI protein product, with the greatest degree of enhancement occurring at the highest doses of rBPI₂₃ and no evidence of the "swamp out" effects observed for combinations with INH intermediate times within the test period.

The results set out above establish utility of BPI protein product concurrently filed in enhancing the growth inhibitory effects of anti-mycobacterial antitotics.

EXAMPLE 6

The following experiment is conducted to determine the *in vitro* inhibitory effect of a BPI protein product on the growth of a *Mycobacterium* species, *Mycobacterium tuberculosis* (MTB). The procedure can be performed with other cultivable *Mycobacterial* species and employs concentrations of a BPI protein product that would be readily generated in human serum by ordinary modes of oral or parenteral administration and/or readily delivered to lung surface by aerosol administration. The effects of the BPI protein product can be evaluated with and without non-ionic surfactants, and/or standard antibiotics.

Log phase cultures of antibiotic-sensitive and antibiotic-resistant MTB are incubated in either 7H11 broth medium or whole human blood, to which the following is added: (a) nothing; (b) surfactant; (c) standard MTB antibiotic; (d) antibiotic plus surfactant. Cultures are incubated with varying concentrations of, e.g., rBPI₂₃. Duplicate cultures grown in each medium are also left untreated by rBPI₂₃ as a negative control. The organisms are placed in Bactec[®] bottles (Johnston Laboratories, Cockeysville, MD) containing ¹⁴C labeled nutrients. rBPI₂₃ challenged *M. tuberculosis* growth is determined by measuring the elution of ¹⁴CO₂ from the medium, compared to the appropriate

negative control. The absence of the formation $^{14}\text{CO}_2$ by the treated cultures is indicative of the inhibitory affects of rBPI₂₃ to MTB. Differential amounts of $^{14}\text{CO}_2$ formed in the absence or presence of standard MTB antibiotics and/or surfactants is indicative of the synergistic or additive effect that a BPI protein product has when used conjunctively with such agents. By comparing the results of this experiment performed with varying concentrations of the BPI protein product, the effective concentration of the BPI protein product is optimized. Radiometric assays to test the susceptibility of *Mycobacterial* species to drugs have been described previously. See McClatchy (cited *supra*) and references therein.

EXAMPLE 7

The following experiment is conducted to determine the *in vitro* effects of a BPI protein product (rBPI₂₃) in an *M. leprae* model. A palmitic acid oxidation assay is used to measure the "viability" of the uncultivable leprosy bacillus adhered to filter paper and "grown" in a ^{14}C -palmitic acid-containing medium. In this method $^{14}\text{CO}_2$ evolved from the metabolism by *M. leprae* of ^{14}C -palmitic acid is trapped on filter paper moistened with NaOH and radioactivity is determined with a liquid scintillation counter. Susceptibility to BPI protein product formulations is determined by differences in radioactivity for *M. leprae* tested with such formulations and treated control cultures.

EXAMPLE 8

The following experiment, which is a variation of an assay conducted by Mittal *et al.*, *J. Clin. Microbiol.*, 17(4):704-707 (1983), is conducted to determine the *in vitro* inhibitory effect of BPI protein product on the growth of *Mycobacterium leprae*. The effects of different concentrations of BPI protein product on *M. leprae* are evaluated with and without non-ionic

surfactants, and/or standard antibiotics. The procedures as described by Mittal *et al.* are outlined below.

Skin biopsy specimens from lepromatous patients are homogenized and are used to inoculate suspensions of mouse peritoneal
5 macrophages cultured in RPMI 1640 (GIBCO Biocult, Irvine, Scotland) enriched with 30% fetal calf serum. After incubating 18 hours, fresh media containing [*methyl*-³H]-thymidine (Amersham International Ltd., Arlington Heights, IL) is added and the cultures are incubated for 14 days. The
10 procedure of Mittal *et al.* is varied by testing the effect of different concentrations of BPI protein product with or without surfactants and/or antibiotics on ³H-thymidine incorporation. Macrophages containing phagocytosed viable *M. leprae* will incorporate ³H-thymidine at a 2 to 10-fold higher rate than control cultures containing heat killed *M. leprae*. Greater than 50% inhibition of ³H-thymidine-incorporation is indicative of bactericidal
15 efficacy of the test product.

EXAMPLE 9

An experiment is conducted to determine the *in vivo* effect that a BPI protein product will have on *M. tuberculosis* species. The model
20 employed is a variation of that used by Lalande *et al.*, *Antimicrobial Agents and Chemotherapy*, 37(3):407-413 (1993), to assess the efficacy of antimicrobial agents against *M. tuberculosis*. Mice inoculated intravenously with *M. tuberculosis* are treated with various BPI protein product doses alone or in combination with surfactants and/or antibiotics. The efficacy of such
25 treatment regimens is analyzed as described.

EXAMPLE 10

The following experiment is conducted to determine the effect that a BPI protein product will have on *M. leprae in vivo*. The model to be
30 used is a variation of that developed by Shepard to study the effect of

compounds on the growth of *M. leprae* in the footpads of infected mice. Shepard *et al.*, *Proc. Soc. Exp. Biol. Med.*, 109:636-638 (1962); Shepard, *J. Exp. Med.* 112:445-454 (1960). Briefly, leprosy bacilli are inoculated into foot-pads of mice, which are subsequently treated with different amounts of test compound with or without known antibiotics and/or surfactants. Untreated infected mice are used as a control. Mice from each treatment regimen are sacrificed at monthly intervals, and sections cut from the infected foot. The presence of an area containing acid-fast bacteria can be observed microscopically and/or the number of such bacteria can be counted. See Shepard and McRae, *Int. J. Lepr.*, 36:78-82. Differences between *M. leprae* bacteria levels observed in treated versus control mice is indicative of the bacteriostatic or bactericidal efficacy of a given BPI treatment regimen. The metabolic status of isolated *M. leprae* may also be measured. Franzblau and Hastings, *Antimicrobial Agents and Chemotherapy*, 31(5):780-783 (1987).

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EXAMPLE 11

The following experiments are designed to demonstrate that BPI protein product is able to inhibit the ability of low concentration of LAM to induce cytokines, yet reverse the unresponsive state that attends higher concentrations of LAM. Increasing concentrations of LAM are pretreated with BPI protein product at varying concentrations. These complexes are applied to peritoneal macrophages from normal and *Mycobacterium* species infected mice. TNF production by treated cells will be assessed.

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EXAMPLE 12

A variation of the armadillo model developed by Kirchheimer *et al.*, *Int. J. Lepr.*, 39:693-702 (1971); *Id.*, 40:229 (1972), is employed to study the *in vivo* effect of BPI protein product test compositions on the growth of *M. leprae* in infected armadillos. Briefly, leprosy bacilli are inoculated into armadillos, which are subsequently treated with different amounts of a test

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composition. The test compositions will comprise a BPI protein product, e.g. rBPI₂₃, with or without known antibiotics and/or surfactants. Untreated infected specimens are used as a control. Armadillos from each treatment regimen are examined and biopsy specimens analyzed by procedures known in the art. *M. leprae* isolated from armadillos is assayed for metabolic activity. Differences between the appearance of lesions, differences in *M. leprae* bacterial concentrations, and differences in the metabolic activity of *M. leprae* isolates in treated versus control specimens are indicative of the bacteriostatic or bactericidal efficacy of a given BPI treatment regimen.

10

EXAMPLE 13

The following experiment is conducted to determine the level of decontamination of a fluid containing LAM that can be achieved by treatment with a BPI protein product. Whole human blood, plasma, blood serum or the like is passed through a column containing a matrix, to which a BPI protein product is bound. Such matrix may be constructed by any means known to those skilled in the art. LAM in the fluid complexes with the BPI protein product affixed to the matrix as the fluid is passed through the column. The absence of LAM in the fluid eluted from the column demonstrates the effectiveness of a BPI protein product at decontaminating a fluid containing LAM.

Alternatively, monoclonal antibodies with binding specificity for a BPI protein product, such as the antibodies employed in Example 1, are affixed to the matrix. A sufficient amount of a BPI protein product is added to the mixture to bind any LAM present in the fluid. The fluid is purified by passing it through the column. The α BPI antibodies affixed to the column bind the LAM/BPI protein product complex in the fluid, and the fluid eluted from the column is analyzed for the presence or absence of LAM contamination.

Numerous modifications and variations in the practice of the invention are expected to occur to those skilled in the art upon consideration of

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the foregoing description of the presently preferred embodiments thereof. For example, while the above illustrative examples principally address studies predictive of antibacterial effects in the context *M.tuberculosis* and *M.leprae*, model studies of infection with, e.g., *M.avium* [see, e.g., Brown *et al.*, 5 *Antimicrob. Agents and Therapy*, 37(3): 398-402(1993)] are also expected to reveal effectiveness of BPI protein product therapies. As another example, preliminary experimental data indicates that BPI protein products alone and/or in combination with cytokines such as gamma interferon (and in combination with antibiotics as well) can enhance the rate at which human monocytes 10- phagocytize Mycobacterial organisms. Combinative therapies involving administration of cytokines along with BPI protein products (and antibiotics) are thus within the scope of the invention). Consequently, the only limitations which should be placed upon the scope of the present invention are those which appear in the appended claims.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: Lewis H. Lambert, Jr.
- (ii) TITLE OF INVENTION: Treatment of Mycobacterial Diseases by Administration of Bactericidal/Permeability-Increasing Protein Product
- (iii) NUMBER OF SEQUENCES: 2
- (iv) CORRESPONDENCE ADDRESS:
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 - (E) COUNTRY: USA
 - (F) ZIP: 60606-6402
- (v) COMPUTER READABLE FORM:
 - (A) MEDIUM TYPE: Floppy disk
 - (B) COMPUTER: IBM PC compatible
 - (C) OPERATING SYSTEM: PC-DOS/MS-DOS
 - (D) SOFTWARE: PatentIn Release #1.0, Version #1.25
- (vi) CURRENT APPLICATION DATA:
 - (A) APPLICATION NUMBER:
 - (B) FILING DATE:
 - (C) CLASSIFICATION:
- (viii) ATTORNEY INFORMATION:
 - (A) NAME: Borun, Michael F.
 - (B) REGISTRATION NUMBER: 25,447
 - (C) REFERENCE/DOCKET NUMBER: 31293
- (ix) TELECOMMUNICATION INFORMATION:
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 - (B) TELEFAX: 312/474-0448
 - (C) TELEX: 25-3856

(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
 - (A) LENGTH: 1813 base pairs
 - (B) TYPE: nucleic acid
 - (C) STRANDEDNESS: single
 - (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: cDNA
- (ix) FEATURE:
 - (A) NAME/KEY: CDS
 - (B) LOCATION: 31..1491
- (ix) FEATURE:
 - (A) NAME/KEY: mat_peptide
 - (B) LOCATION: 124..1491

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

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Gly Thr Ala Val Thr Ala Ala Val Asn Pro Gly Val Val Val Arg Ile	
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Ser Gln Lys Gly Leu Asp Tyr Ala Ser Gln Gln Gly Thr Ala Ala Leu	
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CAG AAG GAG CTG AAG AGG ATC AAG ATT CCT GAC TAC TCA GAC AGC TTT	246
Gln Lys Glu Leu Lys Arg Ile Lys Ile Pro Asp Tyr Ser Asp Ser Phe	
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Lys Ile Lys His Leu Gly Lys Gly His Tyr Ser Phe Tyr Ser Met Asp	
45 50 55	
ATC CGT GAA TTC CAG CTT CCC AGT TCC CAG ATA AGC ATG GTG CCC AAT	342
Ile Arg Glu Phe Gln Leu Pro Ser Ser Gln Ile Ser Met Val Pro Asn	
60 65 70	
GTG GGC CTT AAG TTC TCC ATC AGC AAC GCC AAT ATC AAG ATC AGC GGG	390
Val Gly Leu Lys Phe Ser Ile Ser Asn Ala Asn Ile Lys Ile Ser Gly	
75 80 85	
AAA TGG AAG GCA CAA AAG AGA TTC TTA AAA ATG AGC GGC AAT TTT GAC	438
Lys Trp Lys Ala Gln Lys Arg Phe Leu Lys Met Ser Gly Asn Phe Asp	
90 95 100 105	
CTG AGC ATA GAA GGC ATG TCC ATT TCG GCT GAT CTG AAG CTG GGC AGT	486
Leu Ser Ile Glu Gly Met Ser Ile Ser Ala Asp Leu Lys Leu Gly Ser	
110 115 120	
AAC CCC ACG TCA GGC AAG CCC ACC ATC ACC TGC TCC AGC TGC AGC AGC	534
Asn Pro Thr Ser Gly Lys Pro Thr Ile Thr Cys Ser Ser Cys Ser Ser	
125 130 135	
CAC ATC AAC AGT GTC CAC GTG CAC ATC TCA AAG AGC AAA GTC GGG TGG	582
His Ile Asn Ser Val His Val His Ile Ser Lys Ser Lys Val Gly Trp	
140 145 150	
CTG ATC CAA CTC TTC CAC AAA AAA ATT GAG TCT GCG CTT CGA AAC AAG	630
Leu Ile Gln Leu Phe His Lys Lys Ile Glu Ser Ala Leu Arg Asn Lys	
155 160 165	
ATG AAC AGC CAG GTC TGC GAG AAA GTG ACC AAT TCT GTA TCC TCC AAG	678
Met Asn Ser Gln Val Cys Glu Lys Val Thr Asn Ser Val Ser Ser Lys	
170 175 180 185	
CTG CAA CCT TAT TTC CAG ACT CTG CCA GTA ATG ACC AAA ATA GAT TCT	726
Leu Gln Pro Tyr Phe Gln Thr Leu Pro Val Met Thr Lys Ile Asp Ser	
190 195 200	
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Val Ala Gly Ile Asn Tyr Gly Leu Val Ala Pro Pro Ala Thr Thr Ala	
205 210 215	

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 CATGGTGTGT ATTTTAGGGA TTATGAGCTT CTTTCAAGGG CTAAGGCTGC AGAGATATTT 1731
 CCTCCAGGAA TCGTGTTTCA ATTGTAACCA AGAAATTTCC ATTTGTGCTT CATGAAAAAA 1791
 AACTTCTGGT TTTTTTCATG TG 1813

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 487 amino acids
 (B) TYPE: amino acid
 (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

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 -15 -10 -5 1
 Asn Pro Gly Val Val Val Arg Ile Ser Gln Lys Gly Leu Asp Tyr Ala
 5 10 15
 Ser Gln Gln Gly Thr Ala Ala Leu Gln Lys Glu Leu Lys Arg Ile Lys
 20 25 30
 Ile Pro Asp Tyr Ser Asp Ser Phe Lys Ile Lys His Leu Gly Lys Gly
 35 40 45
 His Tyr Ser Phe Tyr Ser Met Asp Ile Arg Glu Phe Gln Leu Pro Ser
 50 55 60 65
 Ser Gln Ile Ser Met Val Pro Asn Val Gly Leu Lys Phe Ser Ile Ser
 70 75 80
 Asn Ala Asn Ile Lys Ile Ser Gly Lys Trp Lys Ala Gln Lys Arg Phe
 85 90 95
 Leu Lys Met Ser Gly Asn Phe Asp Leu Ser Ile Glu Gly Met Ser Ile
 100 105 110
 Ser Ala Asp Leu Lys Leu Gly Ser Asn Pro Thr Ser Gly Lys Pro Thr
 115 120 125
 Ile Thr Cys Ser Ser Cys Ser Ser His Ile Asn Ser Val His Val His
 130 135 140 145
 Ile Ser Lys Ser Lys Val Gly Trp Leu Ile Gln Leu Phe His Lys Lys
 150 155 160
 Ile Glu Ser Ala Leu Arg Asn Lys Met Asn Ser Gln Val Cys Glu Lys
 165 170 175
 Val Thr Asn Ser Val Ser Ser Lys Leu Gln Pro Tyr Phe Gln Thr Leu
 180 185 190

Pro Val Met Thr Lys Ile Asp Ser Val Ala Gly Ile Asn Tyr Gly Leu
 195 200 205
 Val Ala Pro Pro Ala Thr Thr Ala Glu Thr Leu Asp Val Gln Met Lys
 210 215 220 225
 Gly Glu Phe Tyr Ser Glu Asn His His Asn Pro Pro Pro Phe Ala Pro
 230 235 240
 Pro Val Met Glu Phe Pro Ala Ala His Asp Arg Met Val Tyr Leu Gly
 245 250 255
 Leu Ser Asp Tyr Phe Phe Asn Thr Ala Gly Leu Val Tyr Gln Glu Ala
 260 265 270
 Gly Val Leu Lys Met Thr Leu Arg Asp Asp Met Ile Pro Lys Glu Ser
 275 280 285
 Lys-Phe Arg Leu Thr Thr Lys Phe Phe Gly Thr Phe Leu Pro Glu Val
 290 295 300 305
 Ala Lys Lys Phe Pro Asn Met Lys Ile Gln Ile His Val Ser Ala Ser
 310 315 320
 Thr Pro Pro His Leu Ser Val Gln Pro Thr Gly Leu Thr Phe Tyr Pro
 325 330 335
 Ala Val Asp Val Gln Ala Phe Ala Val Leu Pro Asn Ser Ser Leu Ala
 340 345 350
 Ser Leu Phe Leu Ile Gly Met His Thr Thr Gly Ser Met Glu Val Ser
 355 360 365
 Ala Glu Ser Asn Arg Leu Val Gly Glu Leu Lys Leu Asp Arg Leu Leu
 370 375 380 385
 Leu Glu Leu Lys His Ser Asn Ile Gly Pro Phe Pro Val Glu Leu Leu
 390 395 400
 Gln Asp Ile Met Asn Tyr Ile Val Pro Ile Leu Val Leu Pro Arg Val
 405 410 415
 Asn Glu Lys Leu Gln Lys Gly Phe Pro Leu Pro Thr Pro Ala Arg Val
 420 425 430
 Gln Leu Tyr Asn Val Val Leu Gln Pro His Gln Asn Phe Leu Leu Phe
 435 440 445
 Gly Ala Asp Val Val Tyr Lys
 450 455